Analysis of air filters using the ARL QUANT’X EDXRF Spectrometer

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Introduction
The United States Environmental Protection Agency’s (U.S. EPA) commitment to improve air quality across the USA has created an increasing demand for the monitoring of specific elements in ambient particulate matter collected on filters in residential, industrial and recreational areas. X-ray fluorescence is the preferred method to analyze samples related to air monitoring due to the non-destructive nature of the technique and the superior sensitivity achievable.

The Thermo Scientific™ ARL™ QUANT’X Energy Dispersive X-Ray Fluorescence (EDXRF) instrumentation is ideal for determining concentrations of multiple elements on air filters. The bench top instrument is capable of analyzing the elements sodium to americium with minimal sample preparation and can accommodate filter sizes from 25 mm to 47 mm in diameter using a sample changer or up to 220 mm if loaded manually.

The ARL QUANT’X Spectrometer is equipped with a state-of-the-art silicon drift detector (SDD) which minimizes spectral interference while providing excellent response. Its large active area of 30 mm² ensures a large solid angle for efficient X-ray collection. A high-flux rhodium anode tube, positioned to allow direct excitation from the X-ray tube or tailored excitation through a choice of nine different filters, maximizes sensitivity for a wide range of elements.

Instrument
The key benefits that make the ARL QUANT’X Spectrometer one of the leading instruments in the field of air filter monitoring are as follows:

- Silicon Drift Detector (SDD) with a typical resolution of 135 eV at Mn K-alpha
- Excellent overall sensitivity thanks to superior area of the detector crystal
- High-flux 50 W rhodium anode X-ray tube with excitation voltage of 4 to 50 kV adjustable by steps of 1 kV
- 9-position X-ray tube filter wheel to maximize sensitivity for a wide range of elements
- 10-position automated sample changer (20-position version for 32 mm diameter samples)
- Elemental range sodium (Z=11) through americium (Z=95)
- Multiple element analysis under a single analytical condition
- Wide dynamic range: ppm to percent
- Capable of analyzing various sample types and sizes
- Fast digital pulse processing for increased throughput
- Choice of 5 primary tube collimators (1.0 mm to 8.8 mm)
• Ethernet connectivity for remote access
• Thermo Scientific™ UniQuant™ Software for Standardless XRF Analysis enabling elemental analysis of samples with or without the use of standards
• Thermo Scientific™ WinTrace Software with integrated empirical, fundamental parameters and thin film modules

In addition to the above benefits, the technical features below support an instrument which provides years of reliable service:
• Field transportable and rugged for mobile use
• Low maintenance cost
• Mechanically simple
• Long term stability – individual calibrations remain within QC specifications for months

Excitation conditions
The ARL QUANT’X Spectrometer employs filtered radiation to optimally excite the sample and cause its constituent elements to fluoresce. The ARL QUANT’X Spectrometer is equipped with a nine position filter wheel to reduce spectrum background and tailor excitation energy. The various acquisition parameters are selected for optimal excitation of the elements of concern and best limits of detection.

Sample preparation and presentation
Air filters require no sample preparation prior to analysis except for an inspection of the filter to identify any defects that may affect the analytical results. The filters are analyzed under vacuum conditions to eliminate the ambient atmosphere between the sample and detector thus maximizing light element sensitivity.

The filters are loaded into a 10-position auto-sampler tray if greater than 31 mm in diameter and a 20-position auto-sampler tray if sample is 31 mm diameter or less. The auto-sampler trays feature removable sample holders to accommodate various size filters.

Figure 1: Spectrum of NIST SRM 2783.

Quantitative analysis
Analysis of air filters is accomplished following the U.S. EPA Compendium Method IO-3.3 titled Determination of Metals in Ambient Particulate Matter as modified for use with the ARL QUANT’X Spectrometer. Individual empirical calibrations are achieved using commercial, single and dual element, thin film standards from MicroMatter, Technologies Inc. in addition to blank films. Verification of the calibrations is accomplished by analyzing a National Institute of Standards and Technology (NIST) standard reference material (SRM) 2783 Air particulate on filter media.

Figure 1 shows a spectrum acquired with the ARL QUANT’X Spectrometer on NIST SRM 2783 Air particulate on filter media with elements of interest identified. In this case an excitation voltage of 50 kV, thick Pd filter and a measurement time of 300 s was used. The excellent spectral resolution and associated peak-to-background ratios allows for element lines that are well separated.
Table 1 below shows the five excitation conditions optimized for different groups of elements. The total measurement time per sample is 25 minutes. Measurement time per condition can be further optimized according to specific application needs.

The WinTrace Software uses an advanced deconvolution algorithm to calculate net peak intensities corrected for any overlap by neighboring peaks. This allows for a much more accurate analysis compared to a simple region of interest (ROI) approach around element peaks of interest.

Table 1: Excitation conditions.

<table>
<thead>
<tr>
<th>Condition</th>
<th>Filter</th>
<th>Voltage (kV)</th>
<th>Atmosphere</th>
<th>Live Time (s)</th>
<th>Elements</th>
</tr>
</thead>
<tbody>
<tr>
<td>Low Za</td>
<td>None</td>
<td>4</td>
<td>Vacuum</td>
<td>300</td>
<td>Na, Mg</td>
</tr>
<tr>
<td>Low Zb</td>
<td>Thick C</td>
<td>15</td>
<td>Vacuum</td>
<td>300</td>
<td>Al, Si, P, S, Cl, K, Ca, Sc, Ag, Cd, Sn, Sb, Te, I</td>
</tr>
<tr>
<td>Low Zc</td>
<td>Al</td>
<td>20</td>
<td>Vacuum</td>
<td>300</td>
<td>Ti, V, Cr, Mn, Cs, Ba, La</td>
</tr>
<tr>
<td>Mid Zc</td>
<td>Thick Pd</td>
<td>50</td>
<td>Vacuum</td>
<td>300</td>
<td>Fe, Co, Ni, Cu, Zn, Ga, Ge, As, Se, Br, Rb, Sr, Y, Zr, Hg, Tl, Pb, Bi</td>
</tr>
<tr>
<td>High Za</td>
<td>Thin Cu</td>
<td>50</td>
<td>Vacuum</td>
<td>300</td>
<td>Nb, Mo, Ag, Cd, In, Sn, Sb, Te, I</td>
</tr>
</tbody>
</table>

**Limits of detection**

Minimum detection limits (MDLs) are an important analytical component as they define use limitations of the acquired data. The determination of MDLs for ambient air filter analysis is achieved through the repeated analysis of multiple laboratory blanks. The ARL QUANT’X Spectrometer MDLs displayed in Table 2 are calculated using the average instrument uncertainty from ten laboratory blanks. The ARL QUANT’X Spectrometer data is compared to those specified in the U.S. EPA Compendium Method IO-3.3. The MDLs are reported as $1_{\sigma}$ (68% confidence level) as outlined in the U.S. EPA Compendium Method.

For a majority of elements listed, the MDLs achievable with the ARL QUANT’X Spectrometer are as good as or better than those reported by the U.S. EPA in Compendium Method IO-3.3. The detection levels calculated for the ARL QUANT’X Spectrometer data range from less than 1 ng/cm$^2$ to 21 ng/cm$^2$. For a number of elements listed in Table 2 an extra value is given between parentheses. These are the MDL $1_{\sigma}$ values as calculated for the L lines of these element. Though L lines of these elements typically suffer more from spectral interference than the corresponding K lines, a better MDL is possibly attainable.

A graphical representation of the $1_{\sigma}$ MDLs achievable with the ARL QUANT’X Spectrometer versus those reported by the U.S. EPA in Compendium Method IO-3.3 is presented in Figure 2.
Conclusion
These results show the extraordinary performance that the ARL QUANT’X Spectrometer can achieve for air filter analysis. The detection levels range from less than 1 ng/cm$^2$ to a few ng/cm$^2$. Performance for heavy and light elements alike is extremely good. All ARL QUANT’X Spectrometer detection limits are well below those expected in the EPA’s Compendium Method IO-3.3.